PATENT APPLICATION Docket No.: 20336-00016

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Inventor:

Tariq M. RANA

Application Number:

10/722,176

Filing Date

November 24, 2003

Title

DELIVERY OF siRNAs

Attorney Docket No.:

20336-00016

Examiner

Kimberly Chong

Group Art Unit

1635

Commissioner for Patents P.O. Box 1450 Alexandria, VA 22313-1450

CORRECTED

SECOND DECLARATION OF TARIQ M. RANA UNDER 37 C.F.R. §1.132

I, Tariq M. Rana, hereby make the following declaration:

- 1. I am the inventor of the invention described in U.S. patent application 10/722,176 filed November 24, 2003 ("the '176 application"), and U.S. provisional patent application 60/430,520, filed on November 26, 2002 ("the '520 application").
- 2. The '176 application as filed (and published as US 2004/0204377 A1 on October 12, 2004) describes the use of a delivery mixture comprising a delivery agent consisting of a dendrimer mixed with a nucleic acid capable of mediating RNA interference in Example 1 (paragraph [0102] of the published application and FIGS. 1A and 1B), Example 2 (paragraph [0103] of the published application and FIG. 2), and Example 7 (paragraph [0111] of the published application and FIGS. 9A –9I).
- 3. The corresponding description of the use of a delivery mixture comprising a delivery agent consisting of a dendrimer mixed with a nucleic acid capable of mediating RNA interference can be found in U.S. provisional patent application no. 60/430,520 in Example 2 (page 19 line 23 to page 20, line 31 and FIGS. 1A and 1B), Example 3 (page 21, lines 1-21 and FIG. 2), and Example 8 (page 24, line 29 to page 25, line 14 and FIGS. 9A –9I).

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4. Experiments corresponding to the description of the use of a delivery mixture comprising a delivery agent consisting of a dendrimer mixed with a nucleic acid capable of mediating RNA interference were carried out in my laboratory by Ya-Lin Chiu under my direction and supervision.

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- 5. Exhibits 1-11 are copies of notebook entries from the laboratory notebook of Ya-Lin Chiu, titled "Delivery Method and Localization of siRNA," for the period October 10, 2002 through February 11, 2003. Exhibits 1-11 are notebook entries of experiments carried out during the period October 21, 2002 through November 19, 2002.
 - a. Exhibit 1 is an outline of experimental protocol used in experiments dated 10/21/02 10/25/02 for determination of transfection efficiency of siRNA by various delivery agents, including PAMAM dendrimer and lipofectamine.
 - b. Exhibit 2 is a description of the PAMAM dendrimer used in experiments. A PAMAM G4 dendrimer in a 90 μ g/ μ l methanol solution was used.
 - c. Exhibit 3 is fluorescence level results of siRNA transfection experiments carried out 10/23/02.
 - d. Exhibit 4 is a computer printout of fluorescence results of transfection experiments using PAMAM G4 dendrimer and Lipofectamine, and carried out 10/23/02.
 - e. Exhibit 5 is graph results of transfection experiments introducing Cy3-labeled EGFP duplex siRNAs using PAMAM G4 dendrimer and Lipofectamine carried out 10/23/02. A note in the page indicates the product code number in the Aldrich catalog for the PAMAM Generation 4 dendrimer used in experiments. Exhibit 8 is a graph summary of results of similar transfection experiments introducing Cy3-labeled CDK9 duplex siRNAs carried out 10/23/02.
 - f. Exhibit 6 is protein levels of lysates from cells transfected with CDK9 siRNA by PAMAM dendrimer 10/29/02.

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- g. Exhibit 7 is immunoblot results of lysates from cells transfected with CDK9 siRNA by PAMAM dendrimer. Exhibit 9 is also immunoblot results of lysates from cells transfected with CDK9 siRNA by PAMAM dendrimer on 10/25/02.
- h. Exhibit 10 and Exhibit 11 depict results of microscopic examination of HeLa cells transfected with Cy3-SS/AS siRNA by Lipofectamine or PAMAM, respectively on 11/19/02.
- 6. The notebook pages of Exhibit 2 and Exhibit 5 evidence use of generation 4 PAMAM dendrimer for transfection experiments. Each of the experiments carried out during the period 10/21/02 through 11/19/02 using a PAMAM dendrimer as a delivery agent used generation 4 PAMAM dendrimer.
- 7. Exhibits 1-11 are experiments and results corresponding to Example 1 (paragraph [0102] of the published application and FIGS. 1A and 1B), Example 2 (paragraph [0103] of the published application and FIG. 2), and Example 7 (paragraph [0111] of the published application and FIGS. 9A –9I) of the '176 application.
 - a. Example 1 and FIG. 1A and FIG. 1B correspond to the experiments and results depicted in Exhibits 1, 3, 4, 5, and 8.
 - b. Example 2 and FIG. 2 correspond to the experiments and results in Exhibits 6, 7, and 9.
 - c. Example 7 and FIG. 9 correspond to the experiments and results in Exhibit 11.
- 8. Exhibits 1-11 are experiments and results corresponding to Example 2 (page 19 line 23 to page 20, line 31 and FIGS. 1A and 1B), Example 3 (page 21, lines 1-21 and FIG. 2), and Example 8 (page 24, line 29 to page 25, line 14 and FIGS. 9A –9I) of U.S. provisional patent application no. 60/430,520.
 - a. Example 2 and FIG. 1A and FIG. 1B correspond to the experiments and results depicted in Exhibits 1, 3, 4, 5, and 8.

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- b. Example 3 and FIG. 2 correspond to the experiments and results in Exhibits 6, 7, and 9.
- c. Example 8 and FIG. 9 correspond to the experiments and results in Exhibit 11.
- 9. The experiments described in the '176 application (Example 1 (paragraph [0102] of the published application and FIGS. 1A and 1B), Example 2 (paragraph [0103] of the published application and FIG. 2), and Example 7 (paragraph [0111] of the published application and FIGS. 9A –9I) of the '176 application) used a generation 4 PAMAM dendrimer as a delivery agent mixed with a nucleic acid capable of mediating RNA interference.
- 10. The experiments described in U.S. Provisional application 60/430,520 (Example 2 (page 19 line 23 to page 20, line 31 and FIGS. 1A and 1B), Example 3 (page 21, lines 1-21 and FIG. 2), and Example 8 (page 24, line 29 to page 25, line 14 and FIGS. 9A –9I) of U.S. provisional patent application no. 60/430,520) used a generation 4 PAMAM dendrimer as a delivery agent mixed with a nucleic acid capable of mediating RNA interference.
- 11. I further declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 101 of Title 18 of the United States Code, and that such willful, false statements may jeopardize the validity of the above-identified application or any patent issuing thereon.

Respectfully Submitted,

Date: 11/9/09

Tariq M. Rana

10/21/02 ~ 102402 Determine i rangemen viticiem 43-labeled siRNA as indicator 5'cy3-SS/AS Duplex EGFP Transfection by

1. Lipofectamina 2. Nanoparticle # 1 3. PAMAM Hela cell 6hir incubation at 37°C washed 3x by 5ml PBS Oragen DNA/RNA Extration kit Solution Removed by strong denaturing reagent RNA Small RNA followed by \$ 13000 pm 20 min Isopropanol precipitation 4°c. 13000 pm 30 mis Radisolve perlet in H=D (heat at 90°C 10 min) Detect Cy3 Signal by Flurometer Ya-Li chow 109/110 mL.

90 mg/me.

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EXHIBIT 3

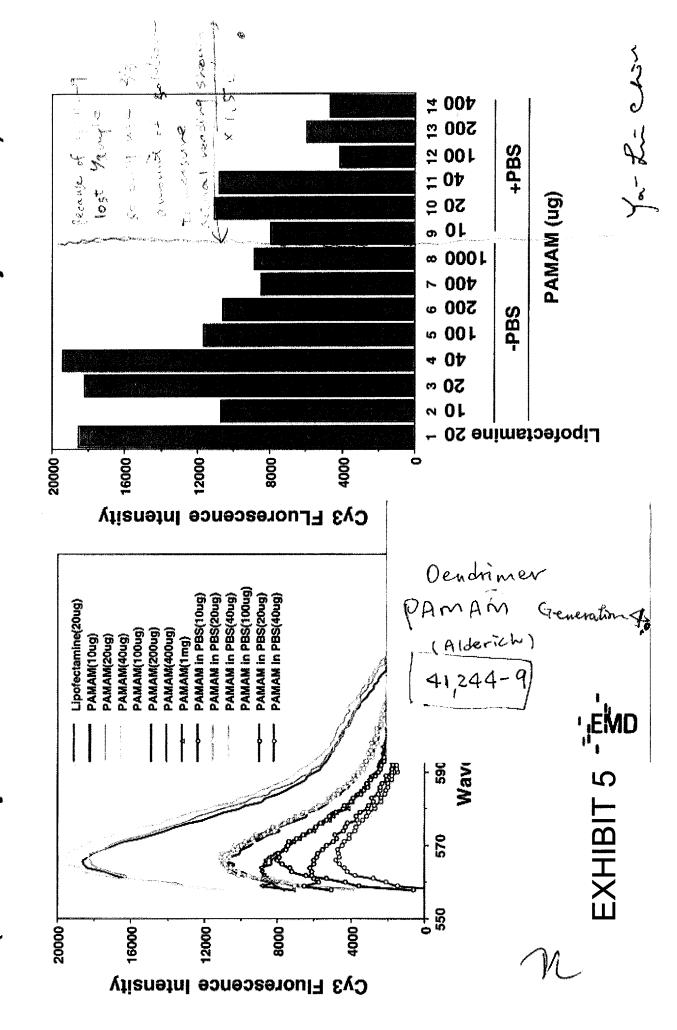
Yo- Richin

102302-1-data2

		Cy3 FLuorescence Intensity	Treatment	Relative siRNA uptake Efficienc
ı	1	18560.000	Lipofectamin (20ug)	1.000
	2	10690.000	PAMAM(10ug)	0.576
	3	18220.000	PAMAM(20ug)	0.982
	4	19400.000	PAMAM(40ug)	1.045
	5	11650.000	PAMAM(100ug)	0.628
	6	10570.000	PAMAM(200ug)	0.570
•	7	8500.000	PAMAM(400ug)	0.458
	8	8867.000	PAMAM(1mg)	0.478
	9	7982.000	PAMAM 1 in PBS (10ug)	0.430
1	0	11080.000	PAMAM 1 in PBS (20ug)	0.597
1	1	10780.000	PAMAM 1 in PBS (40ug)	0.581
1	2	4126.000	PAMAM 1 in PBS (100ug)	0.222
1	3	5962.000	PAMAM 1 in PBS (200ug)	0.321
1	4	4698.000	PAMAM 1 in PBS (400ug)	0.253

Ja-Linek

(EGFP 5'Cy3-SS/AS siRNA as transfection efficiency indicator) Comparison Analysis of PAMAM to Lipofectamine



	OD750	Protein(ug/ul)	60ug	ul/60ug	Buffer	
1 2	0.716 0.613	6.616 5.642	60.000 60.000		11.931 10.366	Mock (young PAMAM only) 1000M cdk9siRNA by 20009 4009
3	0.779	7.211	60.000		12.680	40 mg DAMAM
4	0.783	7.249	60.000	8.277	12.723	100 00
5	0.648	5.973	60.000	10.045	10.955	100 ug PAMAM-C 200 ug 400 ug
6	0.587	5.396	60.000	11.119	9.881	450 00
7	0.553	5.075	60.000	11.823	9,177	1 mg
-8-	0.471	4.300	60,000	13.955	7.045	1
7%		-9-900	60.000	_10 040-	2.000-	

EXHIBIT 6

You to che

20/08/01 ~ 20/55/01

so a cannot use dual fluorescence assay to get Butitation doto but cannot sent the reporter plasmid in (size constraint?) We known that PAMAM Dentrimer can sent sirvid into the cell. Schening of coky expression by pAMAM mediated transfection From 10/21/02 and 10/25/02 beperiment.

Caro ion um coted signifi PAMAM CR Transfected by

EXHIBIT 7

Summar PATITI 40/23/2002 Fig 1 20000 Lipofectamine(20ug) PAMAM(10ug) PAMAM(20ug) Cy3 Fluorescence Intensity 16000 PAMAM(40ug) PAMAM(100ug) PAMAM(200ug) PAMAM(400ug) 12000 PAMAM(1mg) 8000 4000 630 550 570 **590** 610 650 Wavelength(nm) 10/23/2002 1.2 Relative siRNA Uptake Efficiency 1.0 8.0 0.6 0.4 0.2 0.0 2 **0** 20 c 4 04 200 ⁹ 400 1000 [∞] 5 00L PAMAM (ug)

(A)

(B)

EXHIBIT 8

m Ya-Li-ckin Summary of PHMMM

10/25/2002 Fig 2

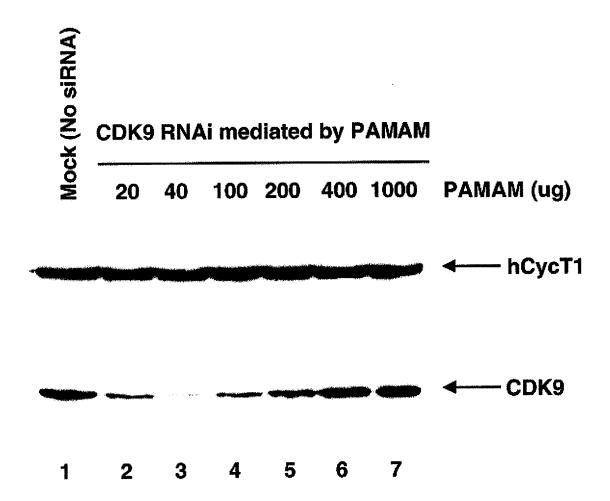
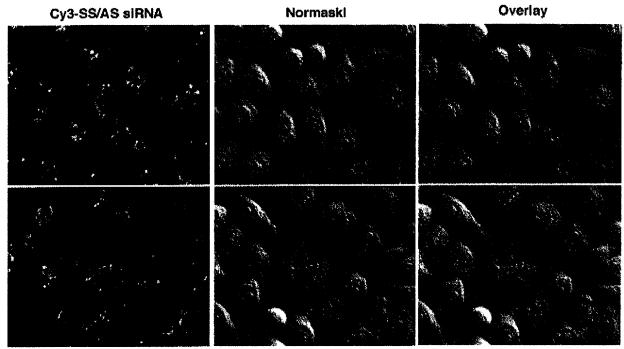


EXHIBIT 9

Ja-Li chin

11/19/02

Uptake of siRNA by HeLa Cells (20ug/1mL Lipofectamine-Mediated Transfection, 6h)(111902)



400X, 35 mm dish with coverslip bottom

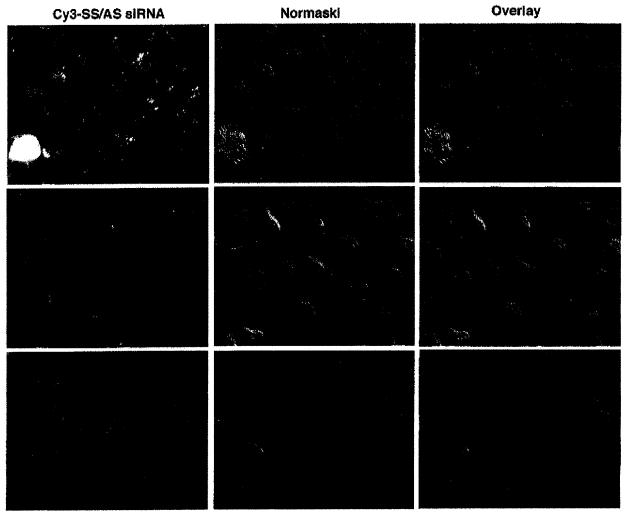
Lelco Demo

EXHIBIT 10

The

1.1. a/2002 Ya- Lin Chie

Uptake of sIRNA by HeLa Cells (40ug/1mL PAMAM-Mediated Transfection, 6h)(111902)



400X, 35 mm dish with coverslip bottom

Leico Demo

EXHIBIT 11

M

W, 912002